## **CLAIMS**

1. Process for the preparation of an enantiomerically enriched compound of formula 1

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$$R^{1}$$
 $C^{*}$ 
 $R^{3}$ 
 $R^{3}$ 
 $R^{4}$ 
 $R^{5}$ 
 $R^{5}$ 

or a salt thereof, wherein:

C\* represents an asymmetric carbon atom;

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R<sup>1</sup> and R<sup>2</sup> are different from each other, and, each independently, represent H, a substituted or unsubstituted alkyl or aryl group;

R<sup>3</sup> represents CH<sub>2</sub>OH or an optionally protected CHO group;

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R<sup>5</sup> represents H, a substituted or unsubstituted alkyl or aryl group; and R<sup>4</sup> represents H or C(=O)R<sup>6</sup> wherein R<sup>6</sup> represents H, a substituted or unsubstituted alkyl, aryl or alkoxy group R<sup>4</sup> represents or an amine protecting group, or R4 and R5 form together with the N to which they are attached a cyclic imide group, wherein an enantiomerically enriched compound with formula 2

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or a salt thereof, wherein  $C^*$ ,  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^5$  are as defined above and  $R^7$  represents H or  $C(=0)R^6$  wherein  $R^6$  is as defined above,  $R^7$  represents an amine protecting group, or  $R^5$  and  $R^7$  form together with the N to which they are attached a cyclic imine group, is subjected to hydrogenation in the presence of hydrogen, a hydrogenation catalyst and a mineral acid.

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2. Process according to claim 1, wherein R<sup>3</sup> is an optionally protected CHO group and wherein hydrogen is present at a hydrogen-pressure between 0.1 and 2 MPa.

- 3. Process according to claim 2, wherein the hydrogen-pressure is between 0.5 and 1 MPa.
- 4. Process according to anyone of claims 1-3 wherein the amino aldehyde is isolated in the form of a chemically and configurationally stable derivative.
- 5 5. Process according to claim 1, wherein R³ is a CH₂OH group and wherein at least during part of the hydrogenation hydrogen is present at a hydrogen-pressure between 2 and 10 MPa.
  - 6. Process according to claim 5, wherein at least during part of the hydrogenation the hydrogen-pressure is between 4 and 6 MPa.
- 7. Process according to claim 5 or 6, wherein the hydrogen-pressure initially is between 0,5 and 2 MPa and subsequently, after most of the nitrile starting material is converted, the hydrogen pressure is increased to a value between 2 and 10 MPa.
- 8. Process according to anyone of claims 1-7 wherein a Pd-catalyst is used as the hydrogenation catalyst.
- Process according to anyone of claims 1-8, wherein as starting material, an enantiomerically enriched nitrile according to formula 2 is used that is prepared by (precursor) fermentation, enzymatic resolution, crystallization induced asymmetric transformation, classical resolution, resolution via preferential crystallization, diastereomeric synthesis, catalytic asymmetric synthesis or dehydratation of amino acid amides.